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	L3	trehalose same L1	263
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=> index bioscience medicine

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, ANTE, AQUALINE, AQUASCI, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CAPLUS, CEABA-VTB, CIN, CONFSCI, CROPB, CROPU, DDFB, DDFU, DGENE, DISSABS, DRUGB, DRUGMONOG2, DRUGU, EMBAL, EMBASE, ...' ENTERED AT 19:36:29 ON 11 JUN 2006

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L1 QUE AMYLASE

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=> file f1-f2, f5-f11, f14, f19

FILE 'CAPLUS' ENTERED AT 19:38:19 ON 11 JUN 2006
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FILE 'AGRICOLA' ENTERED AT 19:38:19 ON 11 JUN 2006

=> s L1

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L3 455 TREHALOSE(S) L2

=> s sulfolobus (s)L3

L4 46 SULFOLOBUS (S) L3

=> dup rem LA

PROCESSING COMPLETED FOR L4

L5 27 DUP REM L4 (19 DUPLICATES REMOVED)

=> s sulfolobales (s)L4

L6 2 SULFOLOBALES (S) L4

=> s L5 (s)L6

L7 2 L5 (S) L6

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L5 ANSWER 1 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2006:113827 USPATFULL << LOGINID::20060611>>

TITLE: Nucleic acid and amino acid sequences relating to Enterobacter cloacae for diagnostics and therapeutics

INVENTOR(S): Weinstock, Keith G., Westborough, MA, UNITED STATES

Deloughery, Craig, Medford, MA, UNITED STATES

Bush, David, Somerville, MA, UNITED STATES

PATENT ASSIGNEE(S): Genome Therapeutics Corporation, Waltham, MA, UNITED STATES (U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 7041814 B1 20060509 APPLICATION INFO.: US 1999-252691 19990218 (9)

NUMBER DATE

PRIORITY INFORMATION: US 1998-94145P 19980724 (60)

US 1998-74787P 19980218 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Smith, Lynette R. F.
ASSISTANT EXAMINER: Portner, Ginny Allen
LEGAL REPRESENTATIVE: Buchanan Ingersoll PC

NUMBER OF CLAIMS: 9 EXEMPLARY CLAIM: 1 LINE COUNT: 19563

AB The invention provides isolated polypeptide and nucleic acid sequences derived from Enterobacter cloacae that are useful in diagnosis and therapy of pathological conditions; antibodies against the polypeptides; and methods for the production of the polypeptides. The invention also provides methods for the detection, prevention and treatment of pathological conditions resulting from bacterial infection.

L5 ANSWER 2 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2005:299054 USPATFULL << LOGINID::20060611>>

Method of producing saccharide preparations TITLE:

INVENTOR(S): Liaw, Gin C., Decatur, IL, UNITED STATES

> Pedersen, Sven, Gentofte, DENMARK Hendriksen, Hanne Vang, Holte, DENMARK Svendsen, Allan, Birkerod, DENMARK Nielsen, Bjarne Ronfeldt, Virum, DENMARK Nielsen, Ruby Illum, Farum, DENMARK

PATENT ASSIGNEE(S): Novozymes A/S, Bagsvaerd, DENMARK (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 2005260719 A1 20051124 APPLICATION INFO.: US 2003-646283 A1 20030821 (10)

RELATED APPLN. INFO.: Continuation of Ser. No. US 2001-908395, filed on 18

Jul 2001, ABANDONED Continuation of Ser. No. US 2000-632392, filed on 4 Aug 2000, GRANTED, Pat. No. US 6303346 Continuation of Ser. No. US 2000-499531, filed on 10 Feb 2000, GRANTED, Pat. No. US 6136571 Continuation of Ser. No. US 1998-198672, filed on 23 Nov 1998, GRANTED, Pat. No. US 6129788 Continuation-in-part of Ser. No. US 1998-107657, filed on 30 Jun 1998, ABANDONED Continuation-in-part of Ser.

No. US 1997-979673, filed on 26 Nov 1997, ABANDONED

DOCUMENT TYPE:

Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: NOVOZYMES NORTH AMERICA, INC., 500 FIFTH AVENUE, SUITE

1600, NEW YORK, NY, 10110, US

NUMBER OF CLAIMS: 11 1-91 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 5 Drawing Page(s)

LINE COUNT:

1205

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method for the production of saccharide preparations, i.e., syrups, by saccharifying a liquefied starch solution, which method comprises a saccharification step during which step one or more enzymatic saccharification stages takes place, and the subsequent steps of one or more high temperature membrane separation steps, and recirculation of the saccharification enzyme, in which method the membrane separation steps are carried out as an integral part of the saccharification step. In another specific aspect, the invention provides a method of producing a saccharide preparation, which method comprises an enzymatic saccharification step, and the subsequent steps of one or more high temperature membrane separation steps and re-circulation of the saccharification enzyme.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 3 OF 27 USPATFULL on STN

2005:4316 USPATFULL << LOGINID::20060611>> ACCESSION NUMBER:

TITLE: Glycosyl hydrolases

INVENTOR(S): Breves, Roland, Mettmann, GERMANY, FEDERAL REPUBLIC OF

Maurer, Karl-Heinz, Erkrath, GERMANY, FEDERAL REPUBLIC

Eck, Jurgen, Heppenheim, GERMANY, FEDERAL REPUBLIC OF Lorenz, Patrick, Lorsch, GERMANY, FEDERAL REPUBLIC OF Zinke, Holger, Zwingenberg, GERMANY, FEDERAL REPUBLIC

NUMBER KIND DATE

PATENT INFORMATION: US 2005003419 Al 20050106 APPLICATION INFO.: US 2004-872874 A1 20040621 (10)

RELATED APPLN. INFO.: Continuation of Ser. No. WO 2002-EP14210, filed on 13 Dec 2002, UNKNOWN

> NUMBER DATE

PRIORITY INFORMATION: DE 2001-10163748 20011221

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: WOODCOCK WASHBURN LLP, ONE LIBERTY PLACE, 46TH FLOOR,

PHILADELPHIA, PA, 19103

NUMBER OF CLAIMS: 21 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 16 Drawing Page(s)

LINE COUNT: 7298

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention relates to a new glycosyl hydrolases with an amyloltic activity and nucleic acids coding for said gylcosyl hydrolases, A PCR-based method for identifying and preparing new gylcosyl hydrolases from metagenome DNA and several possible technical uses for such glycosyl hydrolases with an amylolytic activity. Washing and cleaning products containing such enzymes, and methods and possible uses corresponding thereto are particularly interesting.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 4 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2004:227428 USPATFULL << LOGINID::20060611>>

TITLE: Novel transferase and amylase, process for producing

the enzymes, use thereof, and gene coding for the same

INVENTOR(S): Kato, Masaru, Takasaki-shi, JAPAN

Miura, Yutaka, Takasaki-shi, JAPAN Kettoku, Masako, Takasaki-shi, JAPAN Iwamatsu, Akihiro, Yokohama-shi, JAPAN

Kobayashi, Kazuo, Takasaki-shi, JAPAN Komeda, Toshihiro, Yokohama-shi, JAPAN

PATENT ASSIGNEE(S): KIRIN BEER KUBUSHIKI KAISHA (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 2004175814 A1 20040909 APPLICATION INFO.: US 2003-688276 A1 20031020 (10)

RELATED APPLN. INFO.: Continuation of Ser. No. US 2000-695423, filed on 25

Oct 2000, ABANDONED Continuation of Ser. No. US 1999-298924, filed on 26 Apr 1999, GRANTED, Pat. No. US 6391595 Division of Ser. No. US 1997-750569, filed on 24 Feb 1997, PENDING A 371 of International Ser. No. WO

1995-JP1189, filed on 14 Jun 1995, UNKNOWN

NUMBER DATE

PRIORITY INFORMATION: JP 1994-133354 19940615

JP 1994-194223 19940818 JP 1994-290394 19941031 JP 1994-286917 19941121

JP 1994-311185 19941121

JP 1995-120673 19950421

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: FOLEY AND LARDNER, SUITE 500, 3000 K STREET NW,

WASHINGTON, DC, 20007

NUMBER OF CLAIMS: 145 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 44 Drawing Page(s)

LINE COUNT: 6978

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides a novel transferase that acts on a saccharide, as a substrate, composed of at least three sugar units wherein at least three glucose residues on the reducing end are linked .alpha.-1,4 so as to transfer the .alpha.-1,4 lingages to a .alpha.-1,.alpha.-1 linkages; a process for producing the transferase; a gene coding for the same; and a process for producing an oligosaccharide by using the same. Also provided are a novel amylase that has a principal activity of acting on a saccharide, as a substrate, composed of at least three sugar units wherein at least three sugar units on the reducing end side are glucose units and the linkage between the first and the second glucose units is

.alpha.-1,.alpha.-1 while the linkage between the second and the third glucose units is .alpha.-1,4 so as to liberate .alpha.,.alpha.-trehalose by hydrolyzing the .alpha.-1,4 linkage and another activity of hydrolyzing the .alpha.-1,4 linkage within the molecular chain of the substrate and that liberates disaccharides and/or monosaccharides as the principal final products; a process for producing the amylase; a gene coding for the same; and a process for producing .alpha.,.alpha.-trehalose by using a combination of the transferase and the amylase.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 5 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2002:343879 USPATFULL << LOGINID::20060611>>

TITLE: Novel Polynucleotides

INVENTOR(S): Nakagawa, Satoshi, Tokyo, JAPAN

Mizoguchi, Hiroshi, Tokyo, JAPAN Ando, Seiko, Tokyo, JAPAN Hayashi, Mikiro, Tokyo, JAPAN Ochiai, Keiko, Tokyo, JAPAN Yokoi, Haruhiko, Tokyo, JAPAN Tateishi, Naoko, Tokyo, JAPAN Senoh, Akihiro, Tokyo, JAPAN

Ikeda, Masato, Tokyo, JAPAN

Ozaki, Akio, Hofu-shi, JAPAN

NUMBER KIND DATE

PATENT INFORMATION: US 2002197605 A1 20021226 APPLICATION INFO.: US 2000-738626 A1 20001218 (9)

NUMBER DATE

PRIORITY INFORMATION: JP 1999-377484 19991216

JP 2000-159162 20000407 JP 2000-280988 20000803

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: NIXON & VANDERHYE P.C., 8th Floor, 1100 North Glebe

Road, Arlington, VA, 22201

NUMBER OF CLAIMS: 68 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 4 Drawing Page(s)

LINE COUNT: 13673

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Novel polynucleotides derived from microorganisms belonging to coryneform bacteria and fragments thereof, polypeptides encoded by the polynucleotides and fragments thereof, polynucleotide arrays comprising the polynucleotides and fragments thereof, recording media in which the nucleotide sequences of the polynucleotide and fragments thereof have been recorded which are readable in a computer, and use of them.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

1.5 ANSWER 6 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2002:294680 USPATFULL << LOGINID::20060611>>

TITLE: Method of producing saccharide preparations INVENTOR(S): Liaw, Gin C., Decatur, IL, UNITED STATES

Pedersen, Sven, Gentofte, DENMARK
Hendriksen, Hanne Vang, Holte, DENMARK
Svendsen, Allan, Birkerod, DENMARK
Nielsen, Bjarne Ronfeldt, Virum, DENMARK
Nielsen, Rudy Illum, Farum, DENMARK

PATENT ASSIGNEE(S): Novozymes A/S, Bagsvaerd, DENMARK, DK-2880 (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 2002164723 A1 20021107 APPLICATION INFO.: US 2001-908395 A1 20010718 (9)

RELATED APPLN. INFO.: Continuation of Ser. No. US 2000-632392, filed on 4 Aug

2000, GRANTED, Pat. No. US 6303346 Continuation of Ser. No. US 2000-499531, filed on 10 Feb 2000, GRANTED, Pat. No. US 6136571 Continuation of Ser. No. US 1998-198672, filed on 23 Nov 1998, GRANTED, Pat. No. US 6129788 Continuation-in-part of Ser. No. US 1998-107657, filed on 30 Jun 1998, ABANDONED Continuation-in-part of Ser. No. US 1997-979673, filed on 26 Nov 1997, ABANDONED

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: NOVOZYMES NORTH AMERICA, INC., 500 FIFTH AVENUE, SUITE

1600, NEW YORK, NY, 10110

NUMBER OF CLAIMS: 91 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 5 Drawing Page(s)

LINE COUNT: 1477

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method for the production of saccharide preparations, i.e., syrups, by saccharifying a liquefied starch solution, which method comprises a saccharification step during which step one or more enzymatic saccharification stages takes place, and the subsequent steps of one or more high temperature membrane separation steps, and re-circulation of the saccharification enzyme, in which method the membrane separation steps are carried out as an integral part of the saccharification step.

In another specific aspect, the invention provides a method of producing a saccharide preparation, which method comprises an enzymatic saccharification step, and the subsequent steps of one or more high temperature membrane separation steps and re-circulation of the saccharification enzyme.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 7 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2002:116032 USPATFULL << LOGINID::20060611>>

TITLE: Transferase and amylase, process for producing the

enzymes, use thereof, and gene coding for the same

INVENTOR(S): Kato, Masaru, Takasaki, JAPAN

Miura, Yutaka, Takasaki, JAPAN

PATENT ASSIGNEE(S): Kirin Beer Kabushiki Kaisha, Tokyo, JAPAN (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 6391595 B1 20020521 APPLICATION INFO.: US 1999-298924 19990426 (9) RELATED APPLN. INFO.: Division of Ser. No. US 750569

NUMBER DATE

PRIORITY INFORMATION: JP 1994-133354 19940615

JP 1994-194223 19940818 JP 1994-290394 19941031 JP 1994-286917 19941121 JP 1994-311185 19941121 JP 1995-120673 19950421

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Achutamurthy, Ponnathapu ASSISTANT EXAMINER: Rao, Manjunath N. LEGAL REPRESENTATIVE: Foley & Lardner

NUMBER OF CLAIMS: 38 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 51 Drawing Figure(s); 44 Drawing Page(s)

LINE COUNT: 5088

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides a novel transferase that acts on a saccharide, as a substrate, composed of at least three sugar units wherein at least three glucose residues on the reducing end are linked .alpha.-1,4 so as to transfer the .alpha.-1,4 lingages to a .alpha.-1,.alpha.-1 linkages;

a process for producing the transferase; a gene coding for the same; and a process for producing an oligosaccharide by using the same. Also provided are a novel amylase that has a principal activity of acting on a saccharide, as a substrate, composed of at least three sugar units wherein at least three sugar units on the reducing end side are glucose units and the linkage between the first and the second glucose units is .alpha.-1, alpha.-1 while the linkage between the second and the third glucose units is .alpha.-1,4 so as to liberate .alpha._alpha.-trehalose by hydrolyzing the .alpha.-1,4 linkage and another activity of hydrolyzing the .alpha.-1,4 linkage within the molecular chain of the substrate and that liberates disaccharides and/or monosaccharides as the principal final products; a process for producing the amylase; a gene coding for the same; and a process for producing .alpha._alpha.-trehalose by using a combination of the transferase and the amylase.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 8 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2001:178850 USPATFULL << LOGINID::20060611>>

TITLE: Method of producing saccharide preparations

INVENTOR(S): Liaw, Gin C., Decatur, IL, United States
Pedersen, Sven, Gentofte, Denmark
Hendriksen, Hanne Vang, Holte, Denmark

Svendsen, Allan, Birker.o slashed.d, Denmark Nielsen, Bjarne R.o slashed.nfeldt, Virum, Denmark

Nielsen, Ruby Illum, Farum, Denmark

PATENT ASSIGNEE(S): Novozymes A/S, Bagsvaerd, Denmark (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 6303346 B1 20011016 APPLICATION INFO.: US 2000-632392 20000804 (9)

RELATED APPLN. INFO.: Continuation of Ser. No. US 2000-499531, filed on 10

Feb 2000, now patented, Pat. No. US 6136571 Continuation of Ser. No. US 1998-198672, filed on 23 Nov 1998, now patented, Pat. No. US 6129788 Continuation-in-part of Ser. No. US 1998-107657, filed on 30 Jun 1998, now abandoned Continuation-in-part of Ser. No. US 1997-979673, filed on 26 Nov 1997, now

abandoned DOCUMENT TYPE: Utility

FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Brunsman, David

LEGAL REPRESENTATIVE: Lambiris, Elias J., Garbell, Jason I.

NUMBER OF CLAIMS: 10 EXEMPLARY CLAIM: 1,5,8

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 1032

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method for the production of saccharide preparations, i.e., syrups, by saccharifying a liquefied starch solution, which method comprises a saccharification step during which step one or more enzymatic saccharification stages takes place, and the subsequent steps of one or more high temperature membrane separation steps, and re-circulation of the saccharification enzyme, in which method the membrane separation steps are carried out as an integral part of the saccharification step. In another specific aspect, the invention provides a method of producing a saccharide preparation, which method comprises an enzymatic saccharification step, and the subsequent steps of one or more high temperature membrane separation steps and re-circulation of the saccharification enzyme.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 9 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2000:157207 USPATFULL << LOGINID::20060611>>

TTTLE: Thermostable trehalose-releasing enzyme INVENTOR(S): Ikegami, Shouji, Okayama, Japan Kubota, Michio, Okayama, Japan

Sugimoto, Toshiyuki, Okayama, Japan Miyake, Toshio, Okayama, Japan

PATENT ASSIGNEE(S): Kabushiki Kaisha Hayashibara Seibutsu Kagaku Kenkyujo, Okayama, Japan (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 6150153 20001121 APPLICATION INFO.: US 1997-888158 19970703 (8)

RELATED APPLN. INFO.: Continuation of Ser. No. US 1995-485132, filed on 7 Jun 1995, now patented, Pat. No. US 5723327

NUMBER DATE

PRIORITY INFORMATION: JP 1994-166126 19940625

JP 1995-109130 19950411

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Prats, Francisco LEGAL REPRESENTATIVE: Browdy and Neimark

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 1677

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Disclosed are novel thermostable trehalose-releasing enzyme, and its preparations and uses. The enzyme is obtainable from the culture of microorganisms such as Sulfolobus acidocaldarius (ATCC 33909 and ATCC 49426) and Sulfolobus solfataricus (ATCC 35091 and ATCC 35092), and capable of hydrolyzing at a temperature of over 55.degree. C. the linkage between a trehalose moiety and the remaining glycosyl moiety in a non-reducing saccharide having a trehalose structure as an end unit and having a degree of glucose polymerization of 3 or higher. Trehalose and compositions containing the same are extensively useful in food products, cosmetics and pharmaceuticals.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 10 OF 27 USPATFULL on STN

2000:142138 USPATFULL <<LOGINID::20060611>> ACCESSION NUMBER:

TITLE: Method of producing saccharide preparations Liaw, Gin C., Decatur, IL, United States INVENTOR(S):

Pedersen, Sven, Gentofte, Denmark Hendriksen, Hanne Vang, Holte, Denmark Svendsen, Allan, Birker.o slashed.d, Denmark Nielsen, Bjarne R.o slashed.nfeldt, Virum, Denmark Nielsen, Ruby Illum, Farum, Denmark

PATENT ASSIGNEE(S): Novo Nordisk A/S, Bagsvaerd, Denmark (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 6136571

20001024 APPLICATION INFO.: US 2000-499531 20000210 (9)

RELATED APPLN. INFO.: Continuation of Ser. No. US 1998-198672, filed on 23

Nov 1998 which is a continuation-in-part of Ser. No. US 1998-107657, filed on 30 Jun 1998, now abandoned which is a continuation-in-part of Ser. No. US 1997-979673, filed on 26 Nov 1997, now abandoned

DOCUMENT TYPE: Utility

FILE SEGMENT: Granted

PRIMARY EXAMINER: Brunsman, David

LEGAL REPRESENTATIVE: Zelson, Esq., Steve T., Lambiris, Esq., Elias J.

NUMBER OF CLAIMS: 15 EXEMPLARY CLAIM: 1,8

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 1052

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method for the production of saccharide preparations, i.e., syrups, by saccharifying a liquefied

starch solution, which method comprises a saccharification step during which step one or more enzymatic saccharification stages takes place, and the subsequent steps of one or more high temperature membrane separation steps, and re-circulation of the saccharification enzyme, in which method the membrane separation steps are carried out as an integral part of the saccharification step. In another specific aspect, the invention provides a method of producing a saccharification step, and the subsequent steps of one or more high temperature membrane separation steps and re-circulation of the saccharification enzyme.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 11 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2000:134456 USPATFULL << LOGINID::20060611>>

TITLE: Method of producing saccharide preparations INVENTOR(S): Liaw, Gin C., Decatur, IL, United States

Pedersen, Sven, Gentoste, Denmark Hendriksen, Hanne Vang, Holte, Denmark Svendsen, Allan, Birker.o slashed.d, Denmark Nielsen, Bjarne R.o slashed.nfeldt, Virum, Denmark Nielsen, Ruby Illum, Farum, Denmark

PATENT ASSIGNEE(S): Novo Nordisk A/S, Bagsvaerd, Denmark (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 6129788 20001010 APPLICATION INFO: US 1998-198672 19981123 (9)

RELATED APPLN. INFO.: Continuation of Ser. No. US 1998-107657, filed on 30

Jun 1998, now abandoned which is a continuation of Ser. No. US 1997-979673, filed on 26 Nov 1997, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Brunsman, David

LEGAL REPRESENTATIVE: Zelson, Steve T., Lambiris, Elias J.

NUMBER OF CLAIMS: 16 EXEMPLARY CLAIM: 1,9

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 1248

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method for the production of saccharide preparations, i.e., syrups, by saccharifying a liquefied starch solution, which method comprises a saccharification step during which step one or more enzymatic saccharification stages takes place, and the subsequent steps of one or more high temperature membrane separation steps, and recirculation of the saccharification enzyme, in which method the membrane separation steps are carried out as an integral part of the saccharification step.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 12 OF 27 USPATFULL on STN

ACCESSION NUMBER: 2000:102106 USPATFULL << LOGINID::20060611>>

TITLE: Acid-stable and thermo-stable enzymes derived from

sulfolobus species

INVENTOR(S): Deweer, Philippe, Aalst, Belgium

Amory, Antione, Rixensart, Belgium

PATENT ASSIGNEE(S): Genencor International, Inc., Rochester, NY, United States (U.S. corporation)

19970929 PCT 102(e) date

NUMBER KIND DATE

PATENT INFORMATION: US 6100073 20000808

WO 9602633 19960201

APPLICATION INFO.: US 1997-765939 19970929 (8)

WO 1995-EP2703 19950707 19970929 PCT 371 date

NUMBER DATE

PRIORITY INFORMATION: GB 1994-14224 19940714

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Prats, Francisco

NUMBER OF CLAIMS: 33 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 1507

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Novel acid-stable and thermo-stable enzymes having .alpha.-1,4 hydrolytic activity and a .alpha.-1,6 hydrolytic activity which are derived from strains of the genus Sulfolobus. These enzymes are capable of expressing high levels of .alpha.-1,4 hydrolytic activity, including the maximum .alpha.-1,4 hydrolytic activity thereof, at highly acidic pHs of between about 2.5 and about 4.5. These .alpha.-amylases are further capable of expressing high levels of .alpha.-1,4 hydrolytic activity, including the maximum .alpha.-1,4 hydrolytic activity thereof, at high temperatures of between about 90.degree. C. and about 120.degree. C. Particularly disclosed herein are such enzymes which are derived from strains of the species S. acidocaldarius and, in particular, Sulfolobus acidocaldarius DSM 639. Modified starch degradation (liquefaction and saccharification) processes using these novel enzymes are also disclosed herein.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 13 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 1

ACCESSION NUMBER: 2000:552128 CAPLUS <<LOGINID::20060611>>

DOCUMENT NUMBER: 134:1121

TITLE: Cloning and expression of the gene encoding novel

.alpha.-amylase from Sulfolobus shibatae in

Escherichia coli

AUTHOR(S): Liu, Li; Chen, Wei; Jin, Cheng

CORPORATE SOURCE: Laboratory of Enzymology, Institute of Microbiology,

Chinese Academy of Sciences, Beijing, 100080, Peop.

Rep. China

SOURCE: Weishengwu Xuebao (2000), 40(3), 323-326

CODEN: WSHPA8; ISSN: 0001-6209

PUBLISHER: Kexue Chubanshe DOCUMENT TYPE: Journal LANGUAGE: Chinese

AB A novel .alpha.-amylase gene was amplified from Sulfolobus shibatae by using PCR technique. The amplified 1.7 kb DNA fragment was inserted into an expression vector pBV220 to yield the recombinant plasmid pSBAM. The novel .alpha.-amylase gene in pSBAM was expressed in E. coli. The prodn. of the novel .alpha.-amylase activity reached over 8 units/100 mL of the culture. The mol. wt. of this enzyme was about 61 kD by SDS-PAGE. The expressed novel .alpha.- ***amylase*** protein in E. coli DH5.alpha. accounted for about 20% of the total protein in the recombinant cell; the cooperative action of the novel .alpha.- ***amylase*** and the maltooliigosyltrehalose synthase from ***Sulfolobus*** shibatae was investigated and ***trehalose*** was detected by using HPLC anal. when using amylose and partial starch hydrolyzates as substrates.

L5 ANSWER 14 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:806076 CAPLUS << LOGINID::20060611>>

DOCUMENT NUMBER: 134:159202

TITLE: Novel glycosyltransferase and .alpha.-amylase:

Catalytic mechanism and utilization for trehalose

production

AUTHOR(S): Kato, Masaru; Kobayashi, Kazuo

CORPORATE SOURCE: Applied Research Center, Kirin Brewery Co., Ltd.,

Gunma, 370-1295, Japan

SOURCE: Glycoenzymes (2000), 199-215. Editor(s): Ohnishi,

Masatake. Japan Scientific Societies Press: Tokyo,

Japan.

CODEN: 69AQDK

DOCUMENT TYPE: Conference; General Review

LANGUAGE: English

AB A review with 33 refs. Amylolytic activity that converts sol. starch to .alpha...alpha.-trehalose (trehalose), was found in the cell homogenate of the thermoacidophilic archaeum, Sulfolobus solfataricus KMl. A specially developed novel assay method showed two enzymes to be essential for this activity. The enzymes, a glycosyltransferase and an .alpha.-amylase, were purified to homogeneity and characterized. The glycosyltransferase catalyzed the conversion of maltooligosaccharides to glycosyltrehaloses; the .alpha.-amylase catalyzed the liberation of trehalose from glycosyltrehaloses. The mol. wt. of these enzymes was estd. to be 76 kDa and 61 kDa and the optimum temp. were 70-80.degree.C and 70-85.degree.C, resp. Both had high thermostability. Based on an anal. of the reaction products, and an expt. using 18O-labeled water and 3H-labeled substrates, it was verified that glycosyltransferase transferred an oligomer segment of maltooligosaccharide to the ClOH position of glucose, located at the reducing end of the parental maltooligosaccharide, to produce a glycosyltrehalose with an intramol. reaction. From the observation of intermol. transglycosylation, the catalytic mechanism of glycosyltransferase appears to be essentially a transglycosylation. Anal. of the reaction products, and exptl. findings using 3H-labeled substrates indicated that the .alpha.-amylase hydrolyzed only the .alpha.-1,4 glucosidic linkage adjacent to the trehalose unit of the glycosyltrehaloses. The reactivity of .alpha.-amylase to glycosyltrehaloses was about 15 times greater than that to maltooligosaccharides. Comparison of mol. binding affinities between maltooligosaccharides and glycosyltrehaloses suggested that the subsite affinity at subsite 1 of the .alpha.-amylase (located at the reducing end side) for both substrates was a very important determinant of reactivity. The yield of trehalose from starch was almost 80% using these two enzymes.

The yield of trehalose from starch was almost 80% using these two enzymes.

REFERENCE COUNT: 33 THERE ARE 33 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 15 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 2

ACCESSION NUMBER:

1999:569905 CAPLUS <<LOGINID::20060611>>

DOCUMENT NUMBER:

131:285460

TITLE: High level production of thermostable .alpha.-amylase

from Sulfolobus solfataricus in high-cell density

culture of the food yeast Candida utilis

AUTHOR(S): Miura, Yutaka; Kettoku, Masako; Kato, Masaru;

Kobayashi, Kazuo; Kondo, Keiji

CORPORATE SOURCE: Central Laboratories for Key Technology, Kirin Brewery

Co., Ltd., Yokohama, 236-0004, Japan

SOURCE:

Journal of Molecular Microbiology and Biotechnology

(1999), 1(1), 129-134

CODEN: JMMBFF; ISSN: 1464-1801

PUBLISHER: Horizon Scientific Press

DOCUMENT TYPE:

Journal

LANGUAGE: English

AB The .alpha.- ***amylase*** from ***Sulfolobus*** solfataricus has the com. important ability to hydrolyze glycosyltrehalose and can be used for the prodn. of ***trehalose*** from sol. starch. The authors have produced this enzyme in the food yeast Candida utilis at extremely high levels. Because the S. solfataricus gene was previously shown to be very poorly expressed, the gene was resynthesized based on codons preferentially found in the highly expressed C. utilis glyceraldehyde-3-phosphate dehydrogenase (GAP) gene. Expression of this synthetic gene under the control of the GAP promoter yielded biol. active .alpha.-amylase, accounting for more than 50% of the sol. protein. Comparison of the expression levels of various chimeric constructs of the synthetic and native genes indicated that the prodn. level of the .alpha.-amylase was improved more than 2.times. 104-fold by substituting the native gene with the synthesized one. Northern anal, revealed the formation of short mRNAs in transformants with constructs contg. native gene fragments, suggesting that premature termination of the transcripts is responsible for the low produ. level. The .alpha.-amylase-producing C. utilis cells were grown up to 92 g dry cell wt. per L in a synthetic medium, yielding 12.3 g/l .alpha.-amylase which accounts for up to 27% of total cell proteins.

REFERENCE COUNT: 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 16 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1998:662360 CAPLUS << LOGINID::20060611>>

DOCUMENT NUMBER: 129:241775

Recombinant thermostable enzyme which releases TITLE:

trehalose from non-reducing saccharide

INVENTOR(S): Mitsuzumi, Hitoshi; Kubota, Michio; Sugimoto,

Toshiyuki

PATENT ASSIGNEE(S): Kabushiki Kaisha Hayashibara Seibutsu Kagaku Kenkyujo,

Pat. Specif. (Aust.), 69 pp. SOURCE:

CODEN: ALXXAP DOCUMENT TYPE: Patent

LANGUAGE: English FAMILY ACC. NUM. COUNT: 2 PATENT INFORMATION:

PATENT NO.	KI	ND DATE	Ał	PLICATION	NO. DATE
AU 690285	B2	19980423	AU 19	95-27131	19950721
AU 9527131	Al	19960201			
JP 08336388	A2	19961224	JP 19	95-189760	19950704
JP 3557289	B2	20040825			
US 6027918	Α	20000222	US 19	97-798269	19970211
US 6346394	B1	20020212	US 19	98-55210	19980406
PRIORITY APPLN.	INFO	D.:	JР 1	994-190180	A 19940721
		ЛР 1995-19	0128	A 19950411	
		JP 1995-18	9760	A 19950704	,
		JP 1995-10	9128	A 19950411	
		US 1995-50	5377	A3 1995072	21
		US 1997-79	98269	A1 1997021	11

AB Disclosed is a recombinant thermostable enzyme which has a mol. wt. of about 54,000-64,000 daltons and a pI of about 5.6-6.6, and releases trehalose from non-reducing saccharides having a trehalose structure as an end unit and a degree of glucose polymn. of at least 3. The enzyme has a satisfactorily high thermostability, i.e., it is not substantially inactivated even when incubated in an aq. soln. (pH 7.0) at 85.degree.C for 60 min, and this facilitates the prodn. of trehalose on an industrial scale and in a satisfactorily high yield. Enzyme was purified from recombinant microorganisms expressing the gene for Sulfolobus acidocaldarius ATCC 33909 .alpha.-amylase. This enzyme was used to prep. trehalose-contg. syrup or powder from corn, tapioca or potato starch. The syrup or powder can be used in food, pharmaceuticals and cosmetics.

L5 ANSWER 17 OF 27 USPATFULL on STN

ACCESSION NUMBER: 1998:22090 USPATFULL <<LOGINID::20060611>>

TTTLE: Thermostable trehalose-releasing enzyme, and its

preparation and uses

INVENTOR(S): Ikegami, Shouji, Okayama, Japan

Kubota, Michio, Okayama, Japan Sugimoto, Toshiyuki, Okayama, Japan Miyake, Toshio, Okayama, Japan

PATENT ASSIGNEE(S): Kabushiki Kaisha Hayashibara Seibutsu Kagaku Kenkyujo, Okayama, Japan (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5723327 19980303 APPLICATION INFO.: US 1995-485132 19950607 (8)

> NUMBER DATE

PRIORITY INFORMATION: JP 1994-166126 19940625

JP 1995-109130 19950411

DOCUMENT TYPE: Utility

FILE SEGMENT: Granted

PRIMARY EXAMINER: Lankford, Jr., Leon B. ASSISTANT EXAMINER: Prats, Francisco C. LEGAL REPRESENTATIVE: Browdy and Neimark

NUMBER OF CLAIMS: 15

EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 1775

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Disclosed are novel thermostable trehalose-releasing enzyme, and its preparations and uses. The enzyme is obtainable from the culture of microorganisms such as Sulfolobus acidocaldarius (ATCC 33909 and ATCC 49426) and Sulfolobus solfataricus (ATCC 35091 and ATCC 35092), and capable of hydrolyzing at a temperature of over 55.degree. C. the linkage between a trehalose moiety and the remaining glycosyl moiety in a non-reducing saccharide having a trehalose structure as an end unit and having a degree of glucose polymerization of 3 or higher. Trehalose and compositions containing the same are extensively useful in food products, cosmetics and pharmaceuticals.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 18 OF 27 USPATFULL on STN

ACCESSION NUMBER: 1998:11912 USPATFULL << LOGINID::20060611>>

TITLE:

Thermostable non-reducing saccharide-forming enzyme its

production and uses

INVENTOR(S): Nakada, Tetsuya, Okayama, Japan

Chaen, Hiroto, Okayama, Japan Sugimoto, Toshiyuki, Okayama, Japan Miyake, Toshio, Okayama, Japan

PATENT ASSIGNEE(S): Kabushiki Kaisha Hayashibara Seibutsu Kagaku Kenkyujo,

Okayama, Japan (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 5714368 19980203

APPLICATION INFO.: US 1995-466434 19950606 (8)

NUMBER DATE

PRIORITY INFORMATION: JP 1994-166011 19940624

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Lankford, Jr., Leon B.
ASSISTANT EXAMINER: Prats, Francisco C.
LEGAL REPRESENTATIVE: Browdy and Neimark

NUMBER OF CLAIMS: 23 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 4 Drawing Figure(s); 4 Drawing Page(s)

LINE COUNT: 1534

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Disclosed are novel thermostable non-reducing saccharides-forming enzyme, its preparation and uses. The enzyme is obtainable from the culture of microorganisms such as Sulfolobus acidocaldarius (ATCC 33909 and ATCC 49426) and Sulfolobus solfataricus (ATCC 35091 and ATCC 35092), and capable of forming non-reducing saccharides having a trehalose structure as an end unit when allowed to act on reducing partial starch hydrolysates at a temperature of over 55.degree. C. Glucoamylase and alpha.-glucosidase readily yield trehalose when allowed to act on the non-reducing saccharides. These non-reducing saccharides and trehalose are extensively useful in food products, cosmetics and pharmaceuticals.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 19 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 3

ACCESSION NUMBER: 1997:276

1997:276006 CAPLUS <<LOGINID::20060611>>

DOCUMENT NUMBER: 126:279243
TITLE: Production of trebalose f

Production of trehalose from starch by novel

trehalose-producing enzymes from Sulfolobus

solfataricus KM1

AUTHOR(S): Kobayashi, Kazuo; Komeda, Toshihiro; Miura, Yutaka;

Kettoku, Masako; Kato, Masaru

CORPORATE SOURCE: Applied Bioresearch Cent., Kirin Brewery Co. Ltd.,

Gunma, 370-12, Japan

SOURCE: Journal of Fermentation and Bioengineering (1997),

83(3), 296-298

CODEN: JFBIEX; ISSN: 0922-338X

PUBLISHER: Society for Fermentation and Bioengineering, Japan

DOCUMENT TYPE: Journal LANGUAGE: English

AB A new process for ***trehalose*** (I) prodn. from starch (II) was developed using a novel glycosyltransferase and a novel alpha.
amylase from ***Sulfolobus*** solfataricus KM1. The yield of I from II was 81.5% using the 2 enzymes and a thermostable debranching enzyme. I prodn. was carried out at high temp. (.apprx.60.degree.) and at a high concn. of II with no risk of contamination by microorganisms or retrogradation of II.

L5 ANSWER 20 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1996:209939 CAPLUS <<LOGINID::20060611>>

DOCUMENT NUMBER: 124:252731

TITLE: Sulfolobus acidocaldarius thermostable enzyme forms

trehalose-containing non-reducing saccharide from reducing amylaceous saccharide and recombinant enzyme

use in saccharification and sweetener or syrup

manufacture

INVENTOR(S): Maruta, Kazuhiko; Kubota, Michio; Sugimoto, Toshiyuki PATENT ASSIGNEE(S): Kabushiki Kaisha Hayashibara Seibutsu Kagaku Kenkyujo,

Japan

SOURCE: Can. Pat. Appl., 67 pp.

CODEN: CPXXEB
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE CA 2154307 AA 19960122 CA 1995-2154307 19950720 JP 08084586 А2 19960402 JP 1995-189706 19950704 B2 20040825 JP 3557288 AU 9527132 A1 19960201 AU 1995-27132 19950721 AU 690698 B2 19980430 EP 709461 Al 19960501 EP 1995-305101 19950721 EP 709461 B1 19980930 R: AT, BE, CH, DE, DK, ES, FR, GB, IT, LI, NL, SE AT 171727 E 19981015 AT 1995-305101 19950721 US 5976856 19991102 US 1995-505448 19950721 TW 432112 20010501 TW 1995-84107700 19950725 A 19990713 US 1997-840236 US 5922578 19970411 US 2002102696 A1 20020801 US 1999-419305 19991015 PRIORITY APPLN. INFO.: JP 1994-190183 A 19940721 JP 1995-189706 A 19950704

AB Disclosed is a recombinant thermostable enzyme which has a mol. wt. of about 69,000-79,000 daltons and a pl of about 5.4.-6.4, and forms non-reducing saccharides having a trehalose structure as an end unit from reducing amylaceous saccharides having a degree of glucose polymn. of at least 3. The enzyme has satisfactorily high thermostability, i.e. it is substantially not inactivated even when incubated in an aq. soln. (pH 7.0) at 85.degree.C for 60 min, and this facilitates the prodn. of such non-reducing saccharides on an industrial scale and in a satisfactorily-high yield.

US 1995-505448

L5 ANSWER 21 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 4

A3 19950721

ACCESSION NUMBER: 1996:736480 CAPLUS <<LOGINID::20060611>> DOCUMENT NUMBER: 126:15438

TITLE: Gene cloning and expression of new trehalose-producing

enzymes from the hyperthermophilic archaeum Sulfolobus

solfataricus KM1

AUTHOR(S): Kobayashi, Kazuo; Kato, Masaru; Miura, Yutaka;

Kettoku, Masako; Komeda, Toshihiro; Iwamatsu, Akihiro

CORPORATE SOURCE: Applied Bioresearch Center, Kirin Brewery Co. Ltd.,

Gunma, 370-12, Japan

SOURCE: Bioscience, Biotechnology, and Biochemistry (1996),

60(11), 1882-1885 CODEN: BBBIEJ; ISSN: 0916-8451 Japan Society for Bioscience, Biotechnology, and PUBLISHER: Agrochemistry DOCUMENT TYPE: Journal LANGUAGE: English AB The genes encoding ***trehalose*** -producing enzymes, a glycosyl-***trehalose*** -producing enzyme (glycosyltransferase) and a glycosyltrehalose-hydrolyzing enzyme (.alpha.- ***amylase***), from ***Sulfolobus*** solfataricus KM1 were cloned and expressed in E. coli. The nucleotide sequence of the glycosyltransferase gene and the .alpha.-amylase gene indicated proteins with lengths of 728 and 558 amino acids and mol. masses of 86-kDa and 65 kDa, resp. Regions highly conserved in the .alpha.-amylase family exist in the amino acid sequences of these enzymes. REFERENCE COUNT: 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L5 ANSWER 22 OF 27 PASCAL COPYRIGHT 2006 INIST-CNRS. ALL RIGHTS RESERVED. on STN **DUPLICATE 5** ACCESSION NUMBER: 1996-0406572 PASCAL <<LOGINID::20060611>> COPYRIGHT NOTICE: Copyright .COPYRGT. 1996 INIST-CNRS. All rights reserved. TITLE (IN ENGLISH): Reaction mechanism of a new glycosyltrehalosehydrolyzing enzyme isolated from the hyperthermophilic archaeum, Sulfolobus solfataricus KM1 KATO M.; MIURA Y.; KETTOKU M.; KOMEDA T.; IWAMATSU A.; AUTHOR: KOBAYASHI K. CORPORATE SOURCE: Applied Bioresearch, Kirin Brewery Co., Ltd., 3 Miyaharacho, Takasakishi, Gunma 370-12, Japan SOURCE: Bioscience, biotechnology, and biochemistry, (1996), 60(5), 925-928, 15 refs. ISSN: 0916-8451 DOCUMENT TYPE: Journal BIBLIOGRAPHIC LEVEL: Analytic COUNTRY: Japan LANGUAGE: English AVAILABILITY: INIST-8935, 354000060616350450 AN 1996-0406572 PASCAL <<LOGINID::20060611>> CP Copyright .COPYRGT. 1996 INIST-CNRS. All rights reserved. AB Amylolytic activity, which converts soluble starch to .alpha.,.alpha.-***trehalose*** (***trehalose***), was found in the cell homogenate of the hyperthermophilic acidophilic archaeum, ***Sulfolobus*** solfataricus KM1. Two enzymes, a glycosyltransferase and an .alpha.-***amylase*** , which were essential for this activity were identified. The .alpha.- ***amylase*** was purified to homogeneity on SDS-PAGE.
The .alpha.- ***amylase*** catalyzed the hydrolysis of glycosyltrehaloses to ***trehalose*** . Analysis of the reaction products, kinetic parameters, and experimental findings using .sup.3H-labeled substrates indicated that the .alpha.- ***amylase*** hydrolyzed only the .alpha.-1,4 glucosidic linkage adjacent to the ***trehalose*** unit of the glycosyltrehaloses. Six strains of the Sulfolobaceae family examined were observed to have the glycosyltrehalose-hydrolyzing enzyme, the .alpha.- ***amylase*** . L5 ANSWER 23 OF 27 PASCAL COPYRIGHT 2006 INIST-CNRS. ALL RIGHTS RESERVED. on STN **DUPLICATE 6** ACCESSION NUMBER: 1996-0406409 PASCAL <<LOGINID::20060611>> COPYRIGHT NOTICE: Copyright .COPYRGT. 1996 INIST-CNRS. All rights reserved. TITLE (IN ENGLISH): Reaction mechanism of a new glycosyltrehaloseproducing enzyme isolated from the hyperthermophilic

archaeum, Sulfolobus solfataricus KM1 AUTHOR: KATO M.; MIURA Y.; KETTOKU M.; SHINDO K.; IWAMATSU A.; KOBAYASHI K. CORPORATE SOURCE: Applied Bioresearch Center, Kirin Brewery Co., Ltd., 3 Miyaharacho, Takasakishi, Gunma 370-12, Japan

Bioscience, biotechnology, and biochemistry, (1996),

60(5), 921-924, 15 refs. ISSN: 0916-8451

SOURCE:

DOCUMENT TYPE: Journal BIBLIOGRAPHIC LEVEL: Analytic

COUNTRY:

Japan English

LANGUAGE: AVAILABILITY:

INIST-8935, 354000060616350440 AN 1996-0406409 PASCAL <<LOGINID::20060611>>

CP Copyright .COPYRGT. 1996 INIST-CNRS. All rights reserved.

AB An amylolytic activity, which converts soluble starch to .alpha...alpha.-***trehalose*** (***trehalose***), was found in the cell homogenate of the hyperthermophilic, acidophilic archaeum ***Sulfolobus*** solfataricus KM1. Two enzymes, a glycosyltransferase and an ***amylase***, which are essential for this activity, were purified to homogeneity. A glycosyltransferase catalyzed the conversion of maltooligosaccharides to glycosyltrehaloses. Based on a detailed analysis of the reaction products, kinetic parameters, and an experiment using .sup.3H-labeled substrates, it was verified that glycosyltransferase transferred an oligomer segment of maltooligosaccharide to the Cl-OH position of glucose, located at the reducing end of the maltooligosaccharide, to produce a glycosyltrehalose having an .alpha.-1,1 linkage. The reaction appears to be intramolecular. Nine strains of the Sulfolobaceae family were found to have

L5 ANSWER 24 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 7 ACCESSION NUMBER:

DOCUMENT NUMBER:

glycosyltransferases.

1996:198380 CAPLUS <<LOGINID::20060611>>

124:282714

TITLE:

Purification and characterization of new trehalose-producing enzymes isolated from the hyperthermophilic archae, Sulfolobus solfataricus KM1

AUTHOR(S): Kato, Masaru; Miura, Yutaka; Kettoku, Masako; Shindo,

Kazutoshi; Iwamatsu, Akihiro; Kobayashi, Kazuo

CORPORATE SOURCE: Applied Bioresearch Center, Kirin Brewery Co., Ltd.,

Gunma, 370-12. Japan

SOURCE:

PUBLISHER:

Bioscience, Biotechnology, and Biochemistry (1996),

60(3), 546-50

CODEN: BBBIEJ; ISSN: 0916-8451

Japan Society for Bioscience, Biotechnology, and Agrochemistry

DOCUMENT TYPE: Journal LANGUAGE: English

AB An amylolytic activity that converts sol. starch to .alpha,,.alpha,trehalose (trehalose), was found in the cell homogenate of the hyperthermophilic acidophilic archae, S. solfataricus KM1. DEAE-Toyopearl 650S chromatog. of the homogenate as well as other new reliable assay methods showed 2 enzymes to be essential for this activity. These enzymes, a glycosyltransferase (maltooligosyltrehalose synthase) (I) and an amylase, (maltooligosyltrehalose trehalohydrolase) (II) were purified to homogeneity and characterized. Their mol. wts. were 76 and 61 kDa and activities were maximal at 70-80 and 70-85.degree., resp. High thermostability was noted for each. The reaction products of I and II enzymes on amylooligosaccharides were identified by 1H and 13C NMR spectra and HPLC anal. The cooperative mechanism of the 2 enzymes was used in a new enzymic pathway for trehalose synthesis from starch.

L5 ANSWER 25 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1996:485501 CAPLUS << LOGINID::20060611>>

DOCUMENT NUMBER: 125:188959

Production of trehalose by new trehalose-producing TITLE:

enzymes from the archae

AUTHOR(S): Kobayashi, Kazuo; Kettoku, Masako; Miura, Yutaka;

Kato, Masaru; Komeda, Toshihiro; Iwamatsu, Akihiro

CORPORATE SOURCE: Appl. Bioresearch Cent., Kirin Brew. Co., Ltd.,

Takasaki, 370-12, Japan

SOURCE:

Oyo Toshitsu Kagaku (1996), 43(2), 203-211

CODEN: OTKAE3; ISSN: 1340-3494

PUBLISHER: Nippon Oyo Toshitsu Kagakkai DOCUMENT TYPE: Journal; General Review

LANGUAGE: Japanese

AB A review with 13 refs. on ***trehalose*** prodn. by ***Sulfolobus*** solfataricus, the mechanism of ***trehalose*** formation,

```
***amylase*** , purifn. of these enzymes, physicochem. properties and
  substrate specificities of these enzymes, gene anal. of these enzymes,
  cooperative enzymic prodn. of ***trehalose*** from various substrates,
  and new prodn. method of .alpha.,.alpha. ***trehalose*** from starch.
L5 ANSWER 26 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 8
ACCESSION NUMBER:
                           1996:121151 CAPLUS << LOGINID::20060611>>
DOCUMENT NUMBER:
                            124:169384
TITLE:
                 Cloning and expression of genes for novel transferase
              and amylase of Sulfolobus and uses of the enzymes for
              preparing oligosaccharides
                      Kato, Masaru; Miura, Yutaka; Kettoku, Masako;
INVENTOR(S):
             Iwamatsu, Akihiro; Kobayashi, Kazuo; Komeda, Toshihiro
PATENT ASSIGNEE(S): Kirin Beer K K, Japan
SOURCE:
                   PCT Int. Appl., 357 pp.
              CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                     Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
  PATENT NO.
                     KIND DATE
                                      APPLICATION NO.
                                                             DATE
  WO 9534642
                    A1 19951221 WO 1995-JP1189
                                                         19950614
     W: AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI,
       GB, GE, HU, IS, JP, KE, KG, KR, KZ, LK, LR, LT, LU, LV, MD, MG,
       MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM,
       TT, UA
     RW: KE, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT,
       LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE,
       SN, TD, TG
  AU 9526824
                    Al 19960105 AU 1995-26824
                                                        19950614
                   Al 19970326 EP 1995-921965
  EP 764720
                                                       19950614
  EP 764720
                   B1 20051214
    R: CH, DE, DK, FR, GB, IT, LI
  EP 1130101
                   A2 20010905 EP 2000-125389
                                                       19950614
  EP 1130101
                   A3 20041201
    R: CH, DE, DK, FR, GB, IT, LI
  US 6391595
                   B1 20020521 US 1999-298924
                                                        19990426
  US 2004175814
                     A1 20040909 US 2003-688276
                                                         20031020
PRIORITY APPLN. INFO.:
                                     JP 1994-133354
                                                       A 19940615
                        JP 1994-194223
                                         A 19940818
                        JP 1994-290394
                                         A 19941031
                        JP 1994-286917
                                         A 19941121
                        JP 1994-311185
                                         A 19941121
                        JP 1995-120673
                                         A 19950421
                        EP 1995-921965
                                          A3 19950614
                        WO 1995-JP1189
                                          W 19950614
                        US 1997-750569
                                          A3 19970224
                        US 1999-298924
                                          A1 19990426
                        US 2000-695423
                                          B1 20001025
AB Provided is a novel transferase exhibiting substrate specificity on a
  saccharide (higher than trisaccharide) having .gtoreq.3 glucose residues
  with .alpha.-1,4 linkages at its reducing end. The transferase is able to
  convert the .alpha.-1,4 linkages to .alpha.-1, .alpha.-1 linkages. Also
  provided is a novel amylase exhibiting substrate specificity on a
  saccharide (higher than trisaccharide) having .gtoreq.3 glucose residues,
  which exhibit an .alpha.-1,.alpha.-1 linkage between the 1st and 2nd
  residues and an .alpha.-1,4 linkage between the 2nd and 3rd residues, at
  it reducing end. The amylase is able to produce .alpha.,.alpha.-trehalose
  from the substrates by hydrolyzing the .alpha.-1,4 linkage. The amylase
  is also able to hydrolyze intra-mol. .alpha.-1,4 linkages to produce mono-
  and di-saccharides. The gene encoding the amylase is also isolated and a
  process for producing .alpha.,.alpha.-trehalose using the above described
  transferase and the amylase are also disclosed. The gene encoding the
  transferase and the amylase of both Sulfolobus solfataricus strain KM1 and
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S. acidocaldarius strain ATCC 33909 are isolated and a process for producing oligosaccharides using the enzymes is disclosed.

trehalose -producing enzymes, glycosyltransferase, .alpha.-

L5 ANSWER 27 OF 27 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1992:422222 CAPLUS <<LOGINID::20060611>>

DOCUMENT NUMBER: 117:22222

TITLE: Thermostable amylolytic activity from Sulfolobus

solfataricus Lama, Licia; Nicolaus, Barbara; Trinone, Antonio;

AUTHOR(S):

Morzillo, Pasqualina; Calandrelli, Valeria;

Gambacorta, Agata

CORPORATE SOURCE: Ist. Chim. Mol. Interesse Biol., Arco Felice, Italy

Biotech Forum Europe (1991), 8(4), 201-3 CODEN: BFOEEW; ISSN: 0938-7501 SOURCE:

DOCUMENT TYPE: Journal LANGUAGE: English

AB An amylolytic enzyme (I) from the thermophilic archaebacterium S. solfataricus converted glycogen to glucose and converted sol. starch, amylose, and amylopectin to a mixt. of glucose and trehalose. I was purified 8-fold from the cytoplasmic fraction by successive pptn. with (NH4)2SO4 and Me2CO. I was optimally active at 70.degree. and pH 5.5, and was inhibited by Cu2+ and Zn2+. An intramol. coordinate mechanism appears responsible for trehalose formation.

=> d his

QUE AMYLASE Ll

182875 S L1 L2

- L3 455 S TREHALOSE(S)L2
- 46 S SULFOLOBUS (S)L3 L4
- L5 27 DUP REM L4 (19 DUPLICATES REMOVED)
- 2 S SULFOLOBALES (S)L4 L6
- L7 2 S L5 (S)L6

=> log y